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## WHAT IS CLAIMED IS:

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- 1. An isolated nucleic acid molecule comprising a nucleotide sequence selected from the group consisting of:
- a) the nucleotide sequence as set forth in Figure 1A (SEQ ID NO: 1);
- b) the nucleotide sequence encoding the polypeptide from residues 1-200 or from residues 21-200 as set forth in Figure 1A (SEQ ID NO: 1);
  - c) a nucleotide sequence encoding a polypeptide that is at least about 70 percent identical to the polypeptide as set forth in Figure 1A (SEQ ID NO: 1);
- 15 d) a naturally occurring allelic variant or alternate splice variant of any of (a), (b) or (c);
  - e) a nucleotide sequence complementary to any of (a), (b) or (c);
- f) a nucleotide sequence of (b),(c) or (d) 20 encoding a polypeptide fragment of at least about 25, 50, 75, 100, or greater than 100 amino acid residues;
  - g) a nucleotide sequence of (a), (b) or (c) comprising a fragment of at least about 10, 15, 20, 25, 50, 75, 100, or greater than 100 nucleotides; and
- 25 h) a nucleotide sequence which hybridizes under stringent conditions to any of (a)-(g).
- 2. A $\chi$  isolated nucleic acid molecule comprising a hucleotide sequence selected from the group consisting of: 30
  - a) the nucleotide sequence as set forth in Figure 2A (SEQ \D NO: 11) or Figure 3A (SEQ ID NO: 6) or Figure 12A (SKQ ID NO: 16);
- b) the nucleotide sequence encoding the 35 polypeptide as set forth in Figure 2A (SEO ID NO: 6) from residues 1-322 or from residues 47-322, or as set

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forth in Figure 3A (SEQ ID NO: 11) from residues 1-288 or from residues 19-288, 20-288, 21-288, 22-288, 24-288, or 28-288 or as set forth in Figure 12A from residues 1-302, or from residues 19-302, 20-302, 21-302, 22-302, 24-302 or 28-302;

c) a nucleotide sequence encoding a polypeptide that is at least about 70 percent identical to the polypeptide as set forth in Figure 2A (SEQ ID NO: 6) or Figure 3A (SEQ ID NO: 11) or Figure 12A (SEQ ID NO: 6);

d a naturally occurring allelic variant or alternate splice variant of any of (a), (b) or (c);

- e) a nucleotide sequence complementary to any
  of (a), (b) or (c);
- f) a nucleotide sequence of (b),(c) or (d) encoding a polypeptide fragment of at least about 25, 50, 75, 100, or greater than 100 amino acid residues;
- g) a nucleatide sequence of (a), (b) or (c) comprising a fragment of at least about 10, 15, 20, 25, 20 50, 75, 100, or greater than 100 nucleotides; and h) a nucleotide sequence which hybridizes under stringent conditions to any of (a)-(g).
- 3. The nucleic acid molecule of Claims 1 or 2 wherein the nucleotide sequence is operably linked to an expression control sequence.
  - 4. A host cell comprising the nucleic acid molecule of Claim 2.
  - 5. The host cell of Claim 3 which is a eucaryotic cell.
- 6. The host cell of Claim 3 which is a procaryotic 35 cell.

7. A process for producing a polypeptide comprising growing a culture of the host cell of Claim 3 in suitable culture medium and isolating the polypeptide from the culture.

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- 8. A polypept de produced by the process of Claim
- 9. A polypeptide encoded by the nucleic acid molecule of Claim 1.
  - 10. A polypeptide encoded by the nucleic acid molecule of Claim 2.
  - 11. An isolated polypeptide comprising the amino acid sequence selected from the group consisting of:
    - a) the amino acid sequence as set forth in Figure 1A (SEQ ID NO: 2);
  - b) the mature amino acid sequence as set forth in Figure 1A (SEQ ID NO: 2) comprising a mature amino terminus at residue 21;
    - c) a fragment of the amino acid sequence set forth in Figure 1A (SEQ ID NO: 2) comprising at least about 25, 50, 75, 100, or greater than 100 amino acid residues:
      - d) an ortholog of (a), (b) or (c); and
    - e) an allelic variant or alternative splice variant of (a), (b), (c) or (d).

12. An isolated polypeptide comprising the amino acid sequence selected from the group consisting of:

- a) the amino acid sequence as set forth in Figure 2A (SEQ ID NO: 7) or Figure 3A (SEQ ID NO: 12) or Figure 12A (SEQ ID NO: 17);
- b) the mature amino acid sequence as set forth in Figure 2A (SEQ ID NO: 7) comprising a mature

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amino terminus at residues 47, or Figure 3A (SEQ ID NO: 12) comprising a mature amino terminus at any of residues 19, 20, 21, 22, 24 or 28, or Figure 12A (SEQ ID NO: 17) comprising a mature amino terminus at any of residues 19,20,21,22,24,or 28;

c) a fragment of the amino acid sequence set forth in Figure 2A (SEQ ID NO: 7) or Figure 3A (SEQ ID NO: 12) or Figure 12A (SEQ ID NO: 17) comprising at least about 25, 50, 75, 100, or greater than 100 amino acid residues;

d) an ortholog of (a), (b) or (c); and

- e) an allelic variant or alternative splice variant of (a), (b), (c) or (d).
- 13. An antibody or fragment thereof specifically binding the polypeptide of Claims 9, 10, 11 or 12.
  - 14. The antibody of Claim 11 which is a monoclonal antibody.

15. The antibody of Claim 13 which is a human antibody.

- 16. The antibody of Claim 13 which is a humanized or CDR-grafted antibody.
  - 17. The antibody or fragment of Claim 13 which binds B7RP1 or to a B7RP1 extracellular domain.
- 18. The antibody or fragment of Claim 13 which inhibits the binding of B7RP1 to CRP1.
  - 19. A composition comprising the polypeptide of Claims 9, 10, 11 or 12 and a pharmaceutically acceptable carrier, adjuvant, solubilizer, stabilizer or anti-oxidant.

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20. A polypeptide comprising a derivative of the polypeptide of Claims 9, 10, 11 or 12.

21. The polypeptide of Claim 20 which is covalently modified with a water-soluble polymer.

22. A fusion polypeptide comprising the polypeptide of Claims 9, 10, 11 or 12 fused to a heterologous amino acid sequence.

23. The fusion polypeptide of Claim 22 wherein the heterologous amino acid sequence is an IgG constant domain or fragment thereof.

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24. A method for treating, preventing or ameliorating a T-cell mediated disorder comprising administering to an animal the polypeptide of Claims 9, 10, 11 or 12.

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- 25. A method of diagnosing a T-cell mediated disorder or a susceptibility to a T-cell mediated disorder in an animal comprising:
- a) determining the presence or amount of
   25 expression of the polypeptide of Claims 9, 10, 11 or
   12; and
  - b) diagnosing a T-cell mediated disorder or a susceptibility to a T-cell mediated disorder based on the presence or amount of expression of the polypeptide.
  - 26. A method of identifying a test molecule which binds to a polypeptide comprising:
- a) contacting the polypeptide of Claims 9,35 10, 11 or 12 with a test molecule; and

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- b) determining the extent of binding of the polypeptide to the test molecule.
- 27. The method of Claim 26 further comprising determining the activity of the polypeptide when bound to the compound.
- 28. A method of regulating T-cell activation or proliferation in an animal comprising administering to the animal the nucleic acid molecule of Claims 1, 2 or 3.
  - 29. A transgenic non-human mammal comprising the nucleic acid molecule of Claim 3.
  - 30. A method of suppressing an immune response in an animal comprising administering to the animal an antagonist of CRP1 or B7RP1.
- 31. The method of Claim 30 wherein the antagonist is an antibody which binds B7RP1.
- 32. A method of decreasing IgE production in an animal comprising administering a B7RP1 antagonist or a CRP1 antagonist, or a combination thereof, in an amount effective to decrease IgE production.
  - 33. A method of preventing or treating an IgEmediated disorder comprising administering a therapeutically effective amount of a B7RP1 antagonist, or a CRP1 antagonist, or a combination thereof.
- 34. The method of Claims 32 or 33 wherein the B7RP1 antagonist is an antibody which binds B7RP1 and partially or completely inhibits IgE production.

- 35. The method of Claims 32 or 33 wherein the CRP1 antagonist is an antibody which binds CRP1 and partially or completely inhibits IgE production.
- 5 36. The method of Claim 33 wherein the IgE-mediated disorder is asthma or an allergic disorder.
  - 37. The method of Claims 32 or 33 further comprising administering an IgE antagonist.

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- 38. The method of Claim 37 wherein the IgE antagonist is an anti-IgE antibody.
- 39. A method of enhancing an immune response comprising administering B7RP1, a B7RP1 agonist or a CRP1 agonist.
  - 40. The method of Claim 39 further comprising administering a CD28 agonist.

- 41. The method of claim 39 further comprising administering B7.1 or B7.2 or both.
- 42. A method of treating cancer or viral infection comprising administering B7RP1 or an agonist of CRP1, optionally in combination with B7.1 or B7.2.